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NEWS 1		Web Page for STN Seminar Schedule - N. America
NEWS 2	MAR 31	IFICDB, IFIPAT, and IFIUDB enhanced with new custom IPC display formats
NEWS 3	MAR 31	CAS REGISTRY enhanced with additional experimental spectra
NEWS 4	MAR 31	CA/CAplus and CASREACT patent number format for U.S. applications updated
NEWS 5	MAR 31	LPCI now available as a replacement to LDPCI
NEWS 6	MAR 31	EMBASE, EMBAL, and LEMBASE reloaded with enhancements
NEWS 7	APR 04	STN AnaVist, Version 1, to be discontinued
NEWS 8	APR 15	WPIDS, WPINDEX, and WPIX enhanced with new predefined hit display formats
NEWS 9	APR 28	EMBASE Controlled Term thesaurus enhanced
NEWS 10	APR 28	IMSRESEARCH reloaded with enhancements
NEWS 11	MAY 30	INPAFAMDB now available on STN for patent family searching
NEWS 12	MAY 30	DGENE, PCTGEN, and USGENE enhanced with new homology sequence search option
NEWS 13	JUN 06	EPFULL enhanced with 260,000 English abstracts
NEWS 14	JUN 06	KOREAPAT updated with 41,000 documents
NEWS 15	JUN 13	USPATFULL and USPAT2 updated with 11-character patent numbers for U.S. applications
NEWS 16	JUN 19	CAS REGISTRY includes selected substances from web-based collections
NEWS 17	JUN 25	CA/CAplus and USPAT databases updated with IPC reclassification data
NEWS 18	JUN 30	AEROSPACE enhanced with more than 1 million U.S. patent records

Updated Search

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 NEWS 19  JUN 30  EMBASE, EMBAL, and LEMBASE updated with additional
                  options to display authors and affiliated
                  organizations
 NEWS 20  JUN 30  STN on the Web enhanced with new STN AnaVist
                  Assistant and BLAST plug-in
 NEWS 21  JUN 30  STN AnaVist enhanced with database content from EPFULL
 NEWS 22  JUL 28  CA/CAplus patent coverage enhanced
 NEWS 23  JUL 28  EPFULL enhanced with additional legal status
                  information from the epoline Register
 NEWS 24  JUL 28  IFICDB, IFIPAT, and IFIUDB reloaded with enhancements
 NEWS 25  JUL 28  STN Viewer performance improved
 NEWS 26  AUG 01  INPADOCDB and INPAFAMDB coverage enhanced
 NEWS 27  AUG 13  CA/CAplus enhanced with printed Chemical Abstracts
                  page images from 1967-1998
 NEWS 28  AUG 15  CAOLD to be discontinued on December 31, 2008
 NEWS 29  AUG 15  CAplus currency for Korean patents enhanced

 NEWS EXPRESS JUNE 27 08 CURRENT WINDOWS VERSION IS V8.3,
              AND CURRENT DISCOVER FILE IS DATED 23 JUNE 2008.

 NEWS HOURS    STN Operating Hours Plus Help Desk Availability
 NEWS LOGIN    Welcome Banner and News Items
 NEWS IPC8     For general information regarding STN implementation of IPC 8

Enter NEWS followed by the item number or name to see news on that
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FILE 'HOME' ENTERED AT 18:05:44 ON 20 AUG 2008

=> file reg
COST IN U.S. DOLLARS                                SINCE FILE      TOTAL
                                                         ENTRY    SESSION
FULL ESTIMATED COST                                       0.21       0.21

FILE 'REGISTRY' ENTERED AT 18:05:50 ON 20 AUG 2008
USE IS SUBJECT TO THE TERMS OF YOUR STN CUSTOMER AGREEMENT.
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STRUCTURE FILE UPDATES:   19 AUG 2008  HIGHEST RN 1042061-07-3
DICTIONARY FILE UPDATES:  19 AUG 2008  HIGHEST RN 1042061-07-3

New CAS Information Use Policies, enter HELP USAGETERMS for details.

TSCA INFORMATION NOW CURRENT THROUGH July 5, 2008.
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Updated Search

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L1 STRUCTURE UPLOADED

=> s l1
SAMPLE SEARCH INITIATED 18:09:42 FILE 'REGISTRY'
SAMPLE SCREEN SEARCH COMPLETED - 9617 TO ITERATE

20.8% PROCESSED 2000 ITERATIONS 4 ANSWERS
INCOMPLETE SEARCH (SYSTEM LIMIT EXCEEDED)
SEARCH TIME: 00.00.01

FULL FILE PROJECTIONS: ONLINE **COMPLETE**
BATCH **COMPLETE**
PROJECTED ITERATIONS: 186462 TO 198218
PROJECTED ANSWERS: 121 TO 647

L2 4 SEA SSS SAM L1

=> s l1 full
THE ESTIMATED SEARCH COST FOR FILE 'REGISTRY' IS 177.90 U.S. DOLLARS
DO YOU WANT TO CONTINUE WITH THIS REQUEST? (Y)/N or END:y
FULL SEARCH INITIATED 18:09:46 FILE 'REGISTRY'
FULL SCREEN SEARCH COMPLETED - 194565 TO ITERATE

100.0% PROCESSED 194565 ITERATIONS 415 ANSWERS
SEARCH TIME: 00.00.01

L3 415 SEA SSS FUL L1

=> file hcaplus
COST IN U.S. DOLLARS SINCE FILE ENTRY TOTAL SESSION
FULL ESTIMATED COST 181.12 181.33

FILE 'HCAPLUS' ENTERED AT 18:09:50 ON 20 AUG 2008
USE IS SUBJECT TO THE TERMS OF YOUR STN CUSTOMER AGREEMENT.
PLEASE SEE "HELP USAGETERMS" FOR DETAILS.

FILE COVERS 1907 - 20 Aug 2008 VOL 149 ISS 8
FILE LAST UPDATED: 19 Aug 2008 (20080819/ED)

HCAplus now includes complete International Patent Classification (IPC)
reclassification data for the second quarter of 2008.

New CAS Information Use Policies, enter HELP USAGETERMS for details.

This file contains CAS Registry Numbers for easy and accurate
substance identification.

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=> s l3
L4          271 L3

=> s l3 and diana, g?/au
            271 L3
            162 DIANA, G?/AU
L5            0 L3 AND DIANA, G?/AU

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L7            0 L4 AND YOUNG, D?/AU

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L8            0 L4 AND CHUNDURU, S?/AU

=> s l4 and pd < may 2002
      22731637 PD < MAY 2002
                  (PD<20020500)
L9          245 L4 AND PD < MAY 2002

=> d l9, ibib abs fhitstr, 1-10

L9   ANSWER 1 OF 245  HCAPLUS  COPYRIGHT 2008 ACS on STN
ACCESSION NUMBER:        2002:646274  HCAPLUS
DOCUMENT NUMBER:         137:309665
TITLE:                   Stability of biologically active pyridoxal and
                         pyridoxal phosphate in the presence of lysine
AUTHOR(S):               Huang, Tzou-Chi; Chen, Ming-Hung; Ho, Chi-Tang
CORPORATE SOURCE:        Department of Food Science, National Pingtung
                         University of Science and Technology, Pingtung, 912,
                         Taiwan
SOURCE:                  ACS Symposium Series (2002), 816(Bioactive
                         Compounds in Foods), 143-154
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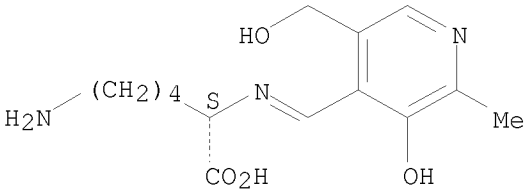
CODEN: ACSMC8; ISSN: 0097-6156
PUBLISHER: American Chemical Society
DOCUMENT TYPE: Journal; General Review
LANGUAGE: English
AB A review on the reactivity of pyridoxal and pyridoxal phosphate toward lysine.
IT 13934-04-8
RL: FMU (Formation, unclassified); FORM (Formation, nonpreparative)
(biol. active pyridoxal and pyridoxal phosphate in presence of lysine)
RN 13934-04-8 HCAPLUS
CN L-Lysine, N2-[[3-hydroxy-5-(hydroxymethyl)-2-methyl-4-pyridinyl]methylene]-
(CA INDEX NAME)

Absolute stereochemistry.
Double bond geometry unknown.



REFERENCE COUNT: 30 THERE ARE 30 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L9 ANSWER 2 OF 245 HCAPLUS COPYRIGHT 2008 ACS on STN
ACCESSION NUMBER: 1999:96142 HCAPLUS
DOCUMENT NUMBER: 130:172994
TITLE: Polymer based pharmaceutical compositions for targeted delivery of biologically active agents
INVENTOR(S): Lau, John R.; Geho, W. Blair
PATENT ASSIGNEE(S): SDG, Inc., USA
SOURCE: PCT Int. Appl., 33 pp.
CODEN: PIXXD2
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 9904824	A1	19990204	WO 1998-US15457	19980724 <--
W: AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ, DE, DK, EE, ES, FI, GB, GE, GH, GM, HR, HU, ID, IL, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, UA, UG, UZ, VN, YU, ZW				
RW: GH, GM, KE, LS, MW, SD, SZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG				
CA 2297025	A1	19990204	CA 1998-2297025	19980724 <--
AU 9885912	A	19990216	AU 1998-85912	19980724 <--
EP 999855	A1	20000517	EP 1998-937127	19980724 <--

Updated Search

R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, MC, PT, IE
JP 2001510811 T 20010807 JP 2000-503875 19980724 <--
PRIORITY APPLN. INFO.: US 1997-53729P P 19970725
WO 1998-US15457 W 19980724

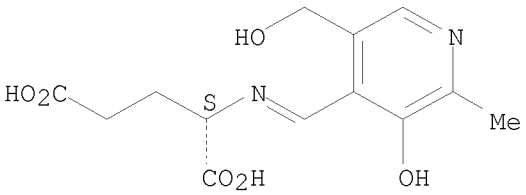
AB A polymeric construct for delivering a biol. active agent to a mammal comprises first polymeric matrix, a biol. active agent contained within the polymeric matrix, and a second polymer chemical bound to the biol. active agent. Said second polymer comprising an amino acid copolymer, said second polymer present in an amount effective to reduce leakage of the active agent from the polymeric construct prior to delivery to the desired situs. A solution contained serotonin HCl (I) 0.07, phytic acid 0.18, polylysine 0.18, polylysine-succinyl 0.18, and N-2,6-(diisopropylphenylacrbamoylmethyl)iminodiacetic acid 0.006 mg/mL. When the solution was filtered through a filter with mol. weight cut-off 3000 about 24.2% of I was retained by the filter, presumably due to ionic and/or hydrogen bonding interaction between I and polymeric component of the solution

IT 13934-03-7
RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses)
(polymer based pharmaceutical compns. for targeted delivery of biol. active agents)

RN 13934-03-7 HCAPLUS

CN L-Glutamic acid, N-[[3-hydroxy-5-(hydroxymethyl)-2-methyl-4-pyridinyl]methylene]- (CA INDEX NAME)

Absolute stereochemistry.
Double bond geometry unknown.



REFERENCE COUNT: 6 THERE ARE 6 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L9 ANSWER 3 OF 245 HCAPLUS COPYRIGHT 2008 ACS on STN
ACCESSION NUMBER: 1999:48605 HCAPLUS
DOCUMENT NUMBER: 130:129967
TITLE: Targeted liposomal constructs for diagnostic and therapeutic uses
INVENTOR(S): Geho, Blair W.; Lau, John R.
PATENT ASSIGNEE(S): SDG, Inc., USA
SOURCE: PCT Int. Appl., 39 pp.
CODEN: PIXXD2
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE

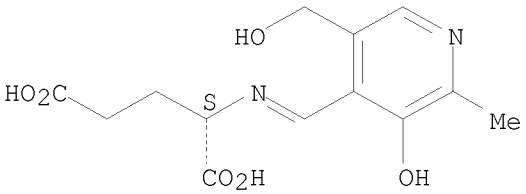
Updated Search

WO 9901110 A1 19990114 WO 1998-US13846 19980702 <--
W: AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ, DE, DK, EE, ES, FI, GB, GE, GH, GM, GW, HR, HU, ID, IL, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, UA, UG, UZ, VN, YU, ZW
RW: GH, GM, KE, LS, MW, SD, SZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, ML, MR, NE, SN, TD, TG
CA 2294900 A1 19990114 CA 1998-2294900 19980702 <--
AU 9882859 A 19990125 AU 1998-82859 19980702 <--
US 6063400 A 20000516 US 1998-109473 19980702 <--
EP 1005327 A1 20000607 EP 1998-933124 19980702 <--
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, MC, PT, IE
JP 2000516641 T 20001212 JP 1999-507412 19980702 <--
MX 9911581 A 20000531 MX 1999-11581 19991213 <--
PRIORITY APPLN. INFO.: US 1997-52740P P 19970702
WO 1998-US13846 W 19980702

AB This invention provides a liposomal construct for delivering a diagnostic or therapeutic agent to a mammal comprising a liposomal carrier, a diagnostic or therapeutic agent entrapped within or associated with the liposomal carrier and a sequestering agent distributed within the liposomal carrier to reduce leakage of the diagnostic or therapeutic agent from the liposomal construct prior to delivery. Claimed liposomal constructs include biogenic amines for deliver them to the hepatocytes. ATP was used as a liposomal sequestrant for serotonin along with the lipid membrane constituents of 1,2-distearoyl-sn-glycerol-3-phosphatidylcholine, dicetyl phosphate, N-(2,6-diisopropylphenylcarbamoylmethyl)iminodiacetic acid and cholesterol.

IT 13934-03-7
RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses)
(targeted liposomal constructs containing diagnostic and therapeutic agents and sequestering agents)
RN 13934-03-7 HCAPLUS
CN L-Glutamic acid, N-[[3-hydroxy-5-(hydroxymethyl)-2-methyl-4-pyridinyl]methylene]- (CA INDEX NAME)

Absolute stereochemistry.
Double bond geometry unknown.



REFERENCE COUNT: 8 THERE ARE 8 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L9 ANSWER 4 OF 245 HCAPLUS COPYRIGHT 2008 ACS on STN
ACCESSION NUMBER: 1999:23873 HCAPLUS
DOCUMENT NUMBER: 130:182286
TITLE: Domain-Structured N1,N2-Derivatized Hydrazines as

10551430

Inhibitors of Ribonucleoside Diphosphate Reductase: Redox-Cycling Considerations

AUTHOR(S): Sarel, Shalom; Fizames, C.; Lavelle, Francois; Avramovici-Grisaru, Shelly

CORPORATE SOURCE: Department of Medicinal Chemistry, Hebrew University of Jerusalem, Jerusalem, 91120, Israel

SOURCE: Journal of Medicinal Chemistry (1999), 42(2), 242-248
CODEN: JMCMAR; ISSN: 0022-2623

PUBLISHER: American Chemical Society

DOCUMENT TYPE: Journal

LANGUAGE: English

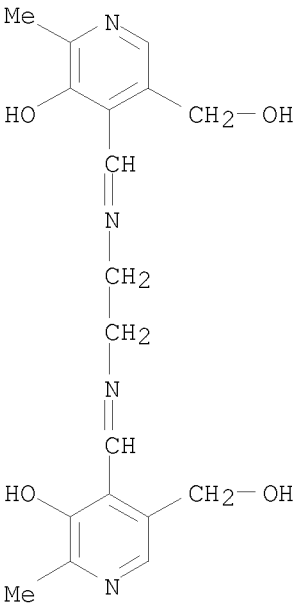
AB Eight analogs of 1-[5-halosalicylidene]-2-[2-pyridinoyl]hydrazine and -[2-pyridyl]hydrazine, four of 1-[pyridoxylidene]-2-[2-pyridinoyl]hydrazine, seven of 1-[pyridoxylidene]-2-[2-pyridyl]hydrazine, and one each of 1,2-bis[pyridoxylidene]diaminoethane and bis[pyridoxylidenehydrazino]phthalazine were synthesized. Their solns. in DMF were assayed for activity against the metalloenzyme ribonucleoside diphosphate reductase (RdR), prepared from a s.c. growing murine tumor (sarcoma 180) implanted in B6D2F3 male mice. The 14C-labeled CDP reductase was assayed by the modified method of Takeda and Weber, in which [14C]cytidine was separated from deoxycytidine by thin-layer chromatog. on cellulose foil. Distribution of radioactivity was assessed with an automatic TLC linear analyzer. Of the 31 compds. tested, 13 were essentially inactive, 7 were highly active against RdR, and the remaining 20 were slightly more active than hydroxyurea (used as a reference compound). The mechanism of inhibition is discussed in terms of three alternative pathways, initiated by sequestration of iron embedded in the R1 subunit of the metalloenzyme to form a C-centered chelate radical (via redox cycling). Alternatively, the latter could either reduce the tyrosyl radical or intercept radicals generated in the reduction process.

IT 88969-07-7P
RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); SPN (Synthetic preparation); BIOL (Biological study); PREP (Preparation)
(preparation and ribonucleoside diphosphate reductase inhibiting activity of pyridinoyl- and pyridylhydrazines)

RN 88969-07-7 HCAPLUS

CN 3-Pyridinemethanol, 4,4'-[1,2-ethanediylbis(nitrilomethylidyne)]bis[5-hydroxy-6-methyl- (CA INDEX NAME)

Updated Search



REFERENCE COUNT: 37 THERE ARE 37 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L9 ANSWER 5 OF 245 HCAPLUS COPYRIGHT 2008 ACS on STN
ACCESSION NUMBER: 1998:458627 HCAPLUS
DOCUMENT NUMBER: 129:241852
ORIGINAL REFERENCE NO.: 129:49163a,49166a
TITLE: Experimental study on a renal imaging agent
AUTHOR(S): Zhu, Jun; Ma, Jixiao; Zhu, Ruisen; Xiong, Jiang; Jin, Changqing
CORPORATE SOURCE: Shanghai 6th People's Hospital, Shanghai, 200233, Peop. Rep. China
SOURCE: Hejishu (1998), 21(5), 297-300
CODEN: NUTEDL; ISSN: 0253-3219
PUBLISHER: Kexue Chubanshe
DOCUMENT TYPE: Journal
LANGUAGE: Chinese
AB The authors report the reactions of glycine, alanine and glycine Et ester with pyridoxal chloride to form the base and the compound chelated with 99mTc in the presence of SnCl2.2H2O. In vivo metabolism was also studied. 99mTc-SB-Gly was rapidly excreted through the kidney into the urine after i.v. injection, with an excretory rate of 79.68±6.66ID% in 30min via urine, a little bit lower than 99mTc-DTPA (82.56±6.88ID%), but having a clear renal scintigraphy. Elimination in blood was rapid. In inhibition expts. with probenecid in rats, the urine excretion rate was not affected, suggesting that this compound passed through by glomerular filtration.
IT 70837-00-2DP, 99mTc complexes
RL: BPR (Biological process); BSU (Biological study, unclassified); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); PROC (Process); USES (Uses)
(biodistribution of renal imaging agents: 99mTc complexes with pyridoxal-amino acid derivs.)
RN 70837-00-2 HCAPLUS

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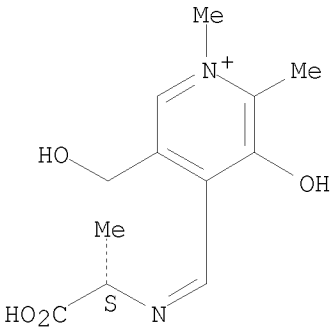
CN Pyridinium, 4-[[[(1S)-1-carboxyethyl]imino]methyl]-3-hydroxy-5-(hydroxymethyl)-1,2-dimethyl-, chloride (9CI) (CA INDEX NAME)

Absolute stereochemistry.
Double bond geometry unknown.



● Cl⁻

L9 ANSWER 6 OF 245 HCAPLUS COPYRIGHT 2008 ACS on STN
ACCESSION NUMBER: 1998:175940 HCAPLUS
DOCUMENT NUMBER: 128:241251
ORIGINAL REFERENCE NO.: 128:47697a,47700a
TITLE: Human salivary proteins CON-1 and CON-2 having alpha-glucosidase-inhibiting activity and their use in treatment of HIV-1 infection and diabetes
INVENTOR(S): Azen, Edwin A.; Pan, David
PATENT ASSIGNEE(S): Wisconsin Alumni Research Foundation, USA
SOURCE: PCT Int. Appl., 55 pp.
CODEN: PIXXD2
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.		DATE
WO 9809981	A1	19980312	WO 1997-US15799		19970908 <--
W: AL, AU, BA, BB, BG, BR, CA, CN, CU, CZ, EE, GE, HU, IL, IS, JP, KP, KR, LC, LK, LR, LT, LV, MG, MK, MN, MX, NO, NZ, PL, RO, SG, SI, SK, TR, TT, UA, UZ, VN, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM					
RW: GH, KE, LS, MW, SD, SZ, UG, ZW, AT, BE, CH, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, ML, MR, NE, SN, TD, TG					
AU 9743359	A	19980326	AU 1997-43359		19970908 <--
US 5981720	A	19991109	US 1997-925237		19970908 <--
PRIORITY APPLN. INFO.:			US 1996-24712P	P	19960909
			WO 1997-US15799	W	19970908

AB Human salivary proteins CON-1 and CON-2 and fragments thereof having alpha-glucosidase inhibitory activity and methods of using same for the

Updated Search

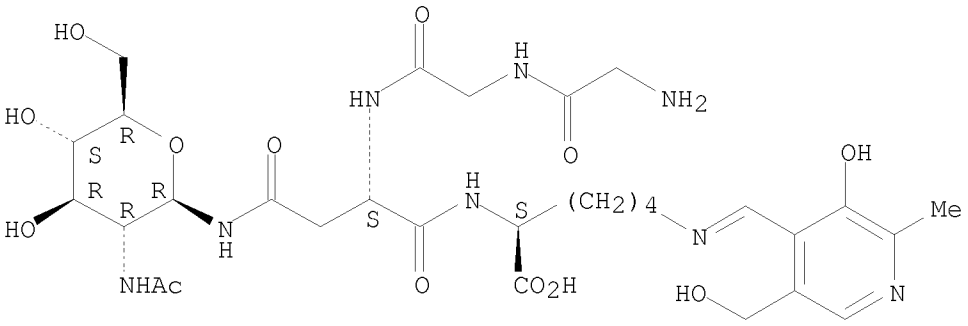
treatment of diabetes and AIDS are disclosed. CON-1 and CON-2 were purified from human saliva. They were found to be glycoproteins. CON-1 inhibited α-glucosidase but removal of carbohydrates from CON-1 decreased its inhibitory activity by 50%. CON-1 reduced HIV-1 proliferation in CEMx174 cells infected with the retrovirus. Protease digestion of CON-1 produced a glycotetrapeptide Gly-Gly-Asn(N-acetyl-β-D-glucosamine)-Lys which also displayed α-glucosidase-inhibiting activity.

IT 204757-17-5
RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); PRP (Properties); THU (Therapeutic use); BIOL (Biological study); USES (Uses)
(human salivary proteins CON-1 and CON-2 having alpha-glucosidase-inhibiting activity and their use in treatment of HIV-1 infection and diabetes)

RN 204757-17-5 HCAPLUS

CN L-Lysine, glycylglycyl-N-[2-(acetylamino)-2-deoxy-β-D-glucopyranosyl]-L-asparaginyl-N6-[[3-hydroxy-5-(hydroxymethyl)-2-methyl-4-pyridinyl]methylene]- (9CI) (CA INDEX NAME)

Absolute stereochemistry.
Double bond geometry unknown.



REFERENCE COUNT: 4 THERE ARE 4 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

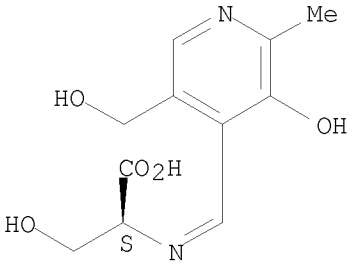
L9 ANSWER 7 OF 245 HCAPLUS COPYRIGHT 2008 ACS on STN
ACCESSION NUMBER: 1998:126640 HCAPLUS
DOCUMENT NUMBER: 128:235002
ORIGINAL REFERENCE NO.: 128:46417a,46420a
TITLE: Skin preparations containing amino acids, antioxidants, and metal-chelating agents
INVENTOR(S): Iwasaki, Keiji; Kitazawa, Manabu
PATENT ASSIGNEE(S): Ajinomoto Co., Inc., Japan
SOURCE: Jpn. Kokai Tokkyo Koho, 22 pp.
CODEN: JKXXAF
DOCUMENT TYPE: Patent
LANGUAGE: Japanese
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE

10551430

JP 10053515 A 19980224 JP 1996-211229 19960809 <--
PRIORITY APPLN. INFO.: JP 1996-211229 19960809
OTHER SOURCE(S): MARPAT 128:235002
AB Skin prepns., which are safe and show long-lasting active O-inhibiting activity, contain ArXCHR(CH2)nY [Ar = (substituted) 2-hydroxyphenyl, 2-hydroxy-1-naphthyl, pyridyl; R = amino acid side chain; X = CH2NH, CH:N; Y = H, CO2R1, SO3H, CONR2R3, CONHCHR5CO2R4; CH2OH; R1-R4 = H, C1-6 alkyl; R5 = amino acid side chain; n = 0, 1] or their salts, antioxidants, and metal-chelating agents. N-(4-pyridoxylmethylene)-L-serine (I), preparation given) 0.1, α-tocopherol 0.5, Na ascorbate 0.5, cetanol 5.0, polyoxyethylene cetyl ether 2.0, olive oil 2.0, propylene glycol 3.0, and H2O to 100 weight% were mixed to give a skin preparation, which was stored at 40° under light irradiation for 3 mo to show 97% I stability.
IT 13933-86-3P
RL: BUU (Biological use, unclassified); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)
(active O-inhibiting skin prepns. containing amino acids, antioxidants, and metal-chelating agents)
RN 13933-86-3 HCAPLUS
CN L-Serine, N-[[3-hydroxy-5-(hydroxymethyl)-2-methyl-4-pyridinyl]methylene]- (CA INDEX NAME)

Absolute stereochemistry.
Double bond geometry unknown.



L9 ANSWER 8 OF 245 HCAPLUS COPYRIGHT 2008 ACS on STN
ACCESSION NUMBER: 1998:126458 HCAPLUS
DOCUMENT NUMBER: 128:205039
ORIGINAL REFERENCE NO.: 128:40559a,40562a
TITLE: Preparation and biological activity of antimicrobial steroidal amino compounds
INVENTOR(S): Schoenecker, Bruno; Wyrwa, Ralf; Moellmann, Ute; Krieg, Reimar; Dubs, Manuela
PATENT ASSIGNEE(S): Friedrich-Schiller-Universitaet Jena, Germany; Hans-Knoell-Institut fuer Naturstofforschung
SOURCE: Ger. Offen., 20 pp.
CODEN: GWXXBX
DOCUMENT TYPE: Patent
LANGUAGE: German
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

PATENT NO. KIND DATE APPLICATION NO. DATE

Updated Search

10551430

DE 19633206	A1	19980219	DE 1996-19633206	19960817 <--
DE 19633206	C2	20010329		
PRIORITY APPLN. INFO.:			DE 1996-19633206	19960817
OTHER SOURCE(S):	MARPAT 128:205039			

GI

* STRUCTURE DIAGRAM TOO LARGE FOR DISPLAY - AVAILABLE VIA OFFLINE PRINT *

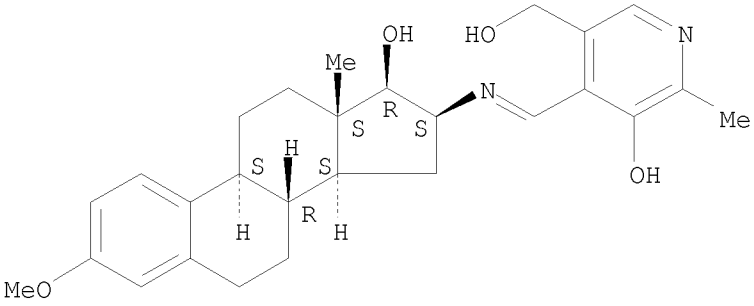
AB Steroidal amines [RNR1R5aCR2R3R4]a+ Aa- [a = 0, 1; R = steroid, cholanyl, cardenolide, bufadienolide derivative; R1 - R5 = H, alkyl; A = anion; when a = 0: R1R2 = bond; R3 = (CH2)xR6, x ≥ 0; R6 = (un)substituted Ph, pyridyl, pyrrolyl, furyl, thienyl, ferrocenyl; R4 = H, alkyl, R3; or when a = 0: R1 = H, alkyl, aryl, acyl,(CH2)yR3, y ≥ 0; R2 = H; R3 = (CH2)xR6; R4 = H, alkyl, R3; when a = 1: R1 = H, alkyl, aryl; R2 = H; R3 = (CH2)xR6; R4 = H, alkyl, R3; R5 = H, alkyl, (CH2)yR7; R7 = (un)substituted Ph, pyridyl, pyrrolyl, furyl, thienyl, ferrocenyl] , [I]a+ Aa- (R8,R9 = H, halo, NO2, OH, alkoxy, aryloxy, acyloxy, acyl, alkyl, aryl; R10 = NR1R5aCR2R3R4), [II]a+ Aa- , [III]a+ Aa- and [IV]a+ Aa- with antimicrobial activity were prepared from the resp. aminosteroids. Steroid I [R1 = R2 = R4 = H, R3 = 2-pyridylmethyl, R8 = β-OH, R9 = OMe, a = 0 (V)] was prepared via reaction of 16β-amino-3-methoxyestra-1,3,5(10)-trien-17β-ol with α-vinylpyridine in MeOH followed by treatment with AcOH. V showed antibacterial activity [25 μg/mL vs. Mycobact. smeg. (SG 987) and Mycobact. fort. B; 12.5 μg/mL vs. Mycobact. chel. B and Mycobact. aurum (SB 66); 12.5 μg/mL vs. Mycobact. vaccae (10670)].

IT 203725-62-6P
RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)
(preparation and antimicrobial activity of steroidal amines)

RN 203725-62-6 HCAPLUS

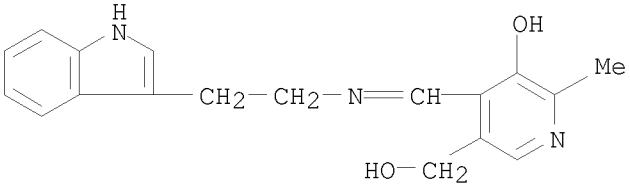
CN Estra-1,3,5(10)-trien-17-ol, 16-[[[3-hydroxy-5-(hydroxymethyl)-2-methyl-4-pyridinyl]methylene]amino]-3-methoxy-, (16β,17β)- (9CI) (CA INDEX NAME)

Absolute stereochemistry.
Double bond geometry unknown.



Updated Search

L9 ANSWER 9 OF 245 HCAPLUS COPYRIGHT 2008 ACS on STN
ACCESSION NUMBER: 1997:294750 HCAPLUS
DOCUMENT NUMBER: 127:41278
ORIGINAL REFERENCE NO.: 127:7783a,7786a
TITLE: Complexes of Mn(II) and Mn(III) with the Schiff base N-[2-(3-ethylindole)]pyridoxaldimine. Electrochemical study of these and related Ni(II) and Cu(II) complexes
AUTHOR(S): Gili, P.; Reyes, M. G. Martin; Zarza, P. Martin; Guedesda Silva, M. F. C.; Tong, Y.-Y.; Pombeiro, A. J. L.
CORPORATE SOURCE: Dep. Quimica Inorganica, Fac. Farmacia, Univ. La Laguna, Tenerife, Canary Islands, Spain
SOURCE: Inorganica Chimica Acta (1997), 255(2), 279-288
CODEN: ICHAA3; ISSN: 0020-1693
PUBLISHER: Elsevier
DOCUMENT TYPE: Journal
LANGUAGE: English
AB New complexes of Mn(II) and Mn(III) with the monoanionic bidentate ligand N-[2-(3-ethylindole)]pyridoxaldimine (pyrdoxTPA) are described. They were characterized by IR and electronic spectroscopies, magnetic measurements and thermogravimetric and calorimetric studies. The spectroscopic and magnetic data indicate a tetrahedral coordination for the Mn(II) complex and a five-coordination for the Mn(III) complex. An electrochem. study of the Mn(II) and analogous Ni(II) and Cu(II) complexes with the same ligand is reported. As indicated by cyclic voltammetry and controlled potential electrolysis, in aprotic medium, the complexes display redox processes involving either the M(II)/M(III) (M = Mn, Ni or Cu) or the M(II)/M(I) (M = Ni or Cu) metal redox pairs, and the pyrdoxTPA ligands. The values of the redox potential of the metal centered redox processes follow the order of those of the corresponding ionization potential of the gaseous metal ions, and for the Mn(II) and Ni(II)complexes evidence is presented for the occurrence of anodically induced trimerizations.
IT 98497-88-2
RL: PRP (Properties); RCT (Reactant); RACT (Reactant or reagent)
(reaction with manganese acetate and elec. potential in DMSO)
RN 98497-88-2 HCAPLUS
CN 3-Pyridinemethanol, 5-hydroxy-4-[[[2-(1H-indol-3-yl)ethyl]imino]methyl]-6-methyl- (CA INDEX NAME)



REFERENCE COUNT: 45 THERE ARE 45 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L9 ANSWER 10 OF 245 HCAPLUS COPYRIGHT 2008 ACS on STN
ACCESSION NUMBER: 1997:244284 HCAPLUS
DOCUMENT NUMBER: 126:232709

10551430

ORIGINAL REFERENCE NO.: 126:44851a,44854a
TITLE: Preparation of magnesium pyridoxal-5'-phosphateglutamate and its intermediate.
INVENTOR(S): Maidonis, Panagiotis; Schneider, Werner
PATENT ASSIGNEE(S): Steigerwald Arzneimittelwerk Gmbh, Germany
SOURCE: Ger. Offen., 8 pp.
CODEN: GWXXBX
DOCUMENT TYPE: Patent
LANGUAGE: German
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
DE 19532625	A1	19970306	DE 1995-19532625	19950904 <--
DE 19532625	C2	20000420		
CA 2230555	A1	19970313	CA 1996-2230555	19960826 <--
WO 9709334	A1	19970313	WO 1996-EP3749	19960826 <--
W: AL, AM, AT, AU, AZ, BB, BG, BR, BY, CA, CH, CN, CU, CZ, DE, DK, EE, ES, FI, GB, GE, HU, IL, IS, JP, KE, KG, KP, KR, KZ, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, TJ, TM, TR, TT, UA, UG, US, UZ, VN				
RW: KE, LS, MW, SD, SZ, UG, AT, BE, CH, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM				
AU 9669845	A	19970327	AU 1996-69845	19960826 <--
AU 706162	B2	19990610		
EP 861258	A1	19980902	EP 1996-930965	19960826 <--
EP 861258	B1	20011121		
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI				
CN 1199402	A	19981118	CN 1996-197528	19960826 <--
HU 9802778	A2	19990928	HU 1998-2778	19960826 <--
HU 9802778	A3	20010228		
JP 11512103	T	19991019	JP 1996-510820	19960826 <--
AT 209210	T	20011215	AT 1996-930965	19960826 <--
ES 2165521	T3	20020316	ES 1996-930965	19960826 <--
PT 861258	T	20020531	PT 1996-930965	19960826
CZ 292662	B6	20031112	CZ 1998-591	19960826
IN 1996CA01512	A	20050304	IN 1996-CA1512	19960826
EG 20974	A	20000830	EG 1996-794	19960831 <--
TW 442472	B	20010623	TW 1996-85110753	19960903 <--
HR 960401	B1	20011031	HR 1996-401	19960903 <--
US 5962687	A	19991005	US 1998-29426	19980629 <--
HK 1014958	A1	20020328	HK 1999-100006	19990104 <--

PRIORITY APPLN. INFO.: DE 1995-19532625 A 19950904
WO 1996-EP3749 W 19960826

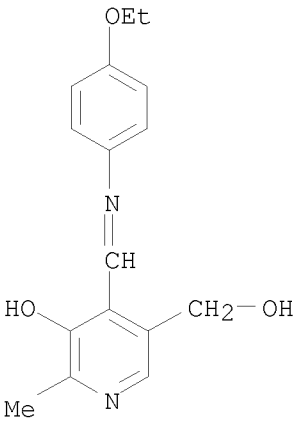
AB Mg5L2 (L5- = pyridoxal-5'-phosphateglutamate) was prepared by the reaction of Mg glutamate and pyridoxal-5'-phosphate. Pyridoxal-5'-phosphate was prepared by a stepwise method starting from pyridoxin hydrochloride oxidation by MnO_2 giving pyridoxal which was reacted with p-phenetidine. P-phenetidylpyridoxal was prepared by this latter reaction and reacted with polyphosphoric acid to give p-phenetidylpyridoxal-5'-phosphate which was deprotected to give pyridoxal-5'-phosphate.

IT 4943-90-2P
RL: IMF (Industrial manufacture); RCT (Reactant); PREP (Preparation); RACT (Reactant or reagent)

Updated Search

10551430

(for preparation of magnesium pyridoxal-5'-phosphateglutamate)
RN 4943-90-2 HCAPLUS
CN 3-Pyridinemethanol, 4-[[(4-ethoxyphenyl)imino]methyl]-5-hydroxy-6-methyl-
(CA INDEX NAME)



=> d his

(FILE 'HOME' ENTERED AT 18:05:44 ON 20 AUG 2008)

FILE 'REGISTRY' ENTERED AT 18:05:50 ON 20 AUG 2008
L1 STRUCTURE UPLOADED
L2 4 S L1
L3 415 S L1 FULL

FILE 'HCAPLUS' ENTERED AT 18:09:50 ON 20 AUG 2008
L4 271 S L3
L5 0 S L3 AND DIANA, G?/AU
L6 0 S L4 AND BAILEY, T?/AU
L7 0 S L4 AND YOUNG, D?/AU
L8 0 S L4 AND CHUNDURU, S?/AU
L9 245 S L4 AND PD < MAY 2002

=> file caold

COST IN U.S. DOLLARS	SINCE FILE ENTRY	TOTAL SESSION
FULL ESTIMATED COST	135.20	316.53

DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS)	SINCE FILE ENTRY	TOTAL SESSION
CA SUBSCRIBER PRICE	-8.00	-8.00

FILE 'CAOLD' ENTERED AT 18:27:51 ON 20 AUG 2008

Updated Search

10551430

FILE COVERS 1907-1966
FILE LAST UPDATED: 01 May 1997 (19970501/UP)

This file contains CAS Registry Numbers for easy and accurate substance identification. Title keywords, authors, patent assignees, and patent information, e.g., patent numbers, are now searchable from 1907-1966. TIFF images of CA abstracts printed between 1907-1966 are available in the PAGE display formats.

New CAS Information Use Policies, enter HELP USAGETERMS for details.

This file supports REG1stRY for direct browsing and searching of all substance data from the REGISTRY file. Enter HELP FIRST for more information.

=> s l3
L10 16 L3

=> d l10, all, 1-16

L10 ANSWER 1 OF 16 CAOLD COPYRIGHT 2008 ACS on STN
AN CA65:18644e CAOLD
TI conversion of 10β, 17β-dehydroxyestra-1,4-dien-3-one to 3-aminoestra-1,3,5(10)-trien-17β-ol
AU Schmialek, Peter; Danneberg, H.
IT 549-02-0 10427-24-4 13144-83-7

L10 ANSWER 2 OF 16 CAOLD COPYRIGHT 2008 ACS on STN
AN CA65:12665f CAOLD
TI formation of pyridoxal phosphate Schiff's base-inherent defect in the tryptophan load test
AU Hughes, P. A. M.; Raine, D. N.
IT 59-00-7 13311-34-7 13311-40-5

L10 ANSWER 3 OF 16 CAOLD COPYRIGHT 2008 ACS on STN
AN CA64:8154c CAOLD
TI pyridine derivs. (S-containing)
PA Merck, E., A.-G.
DT Patent

PATENT NO.	KIND	DATE
NL 6412891		
BE 655454		
GB 1032377		

IT 4632-27-3 4943-89-9 4943-90-2
4943-91-3 4943-92-4 4943-93-5 4943-94-6 4943-95-7
4943-96-8 4943-97-9 4943-98-0 4943-99-1 4944-00-7 4944-01-8
4944-02-9 4944-03-0 4944-04-1 4959-62-0 4959-63-1 4959-64-2
4959-65-3 4959-66-4 4959-67-5 4999-97-7 4999-98-8 4999-99-9
5000-00-0 5000-01-1 5000-02-2 5000-03-3 5000-04-4 5000-05-5
5000-06-6 5000-07-7 5000-08-8 5000-09-9 5000-10-2 5000-11-3
5000-12-4 5000-13-5 5000-14-6 5004-89-7 5004-90-0 5009-62-1
5196-15-6 5196-16-7 5196-17-8 5196-18-9 5365-50-4 5365-58-2
5365-59-3 5365-60-6 5365-61-7 5365-62-8 5365-63-9 5365-64-0
5365-65-1 5365-66-2 5508-97-4 5508-98-5 5572-76-9 5575-18-8

Updated Search

5589-33-3 5589-34-4 30587-24-7 30587-25-8 30587-26-9 30644-49-6
91252-36-7 106504-00-1

L10 ANSWER 4 OF 16 CAOLD COPYRIGHT 2008 ACS on STN
AN CA64:8154b CAOLD
TI pyridoxal Schiff bases
AU Murakami, Masuo; Iwanami, M.; Kawai, R.
PA Yamanouchi Pharmaceutical Co., Ltd.
DT Patent

PATENT NO.	KIND	DATE
JP 65026820		1965

IT 4943-87-7 4943-88-8 5004-88-6

L10 ANSWER 5 OF 16 CAOLD COPYRIGHT 2008 ACS on STN
AN CA63:19f CAOLD
TI reaction of pyridoxal phosphate with amines and its anal. application
AU Gaudiano, Aldo; Polizzi-Sciarrone, M.
IT 54-47-7 66-72-8 1499-44-1 1499-45-2

L10 ANSWER 6 OF 16 CAOLD COPYRIGHT 2008 ACS on STN
AN CA62:12048b CAOLD
TI anomalous rotatory dispersion of metal chelates of aldimines of α-amino acids and their derivs.-determination of absolute configuration
AU Torchinskii, Yu. M.; Koreneva, L. G.
IT 2949-29-3 3269-00-9 3269-01-0 3269-02-1 3444-19-7
3444-20-0 3444-21-1 3444-22-2 3444-23-3 3444-24-4 3444-25-5
3444-26-6 3444-27-7 3444-28-8 3444-29-9 3487-08-9
3520-81-8 3577-08-0 3908-17-6 4055-44-1

L10 ANSWER 7 OF 16 CAOLD COPYRIGHT 2008 ACS on STN
AN CA57:15481a CAOLD
TI erythropoietin
AU De Ritis, Giancarlo
TI semicarbazone formation from pyridoxal, pyridoxal phosphate, and their Schiff bases
AU Cordes, Eugene H.; Jencks, W. P.
IT 781-66-8 1499-44-1 76532-72-4 91761-12-5
93353-85-6 93606-21-4 93688-51-8 96218-00-7

L10 ANSWER 8 OF 16 CAOLD COPYRIGHT 2008 ACS on STN
AN CA56:9354c CAOLD
TI effects of various hormones on the activity and systemic content of histaminase
AU Negishi, Tadamichi
IT 125-04-2 302-25-0 979-32-8 6151-12-8 13331-81-2
13331-82-3 17433-39-5 73622-67-0 73713-65-2 73758-58-4
73840-48-9 73840-49-0 73840-50-3 74037-54-0 82276-93-5
91982-30-8 93884-10-7

L10 ANSWER 9 OF 16 CAOLD COPYRIGHT 2008 ACS on STN
AN CA56:1698c CAOLD
TI chelation therapy in circulatory and sclerosing diseases
AU Boyle, Albert J.; Clarke, N. E.; Mosher, R. E.; McCann, D. S.
TI metal-binding by pyridoxal derivs. and possible relations to tryptophan metabolism

10551430

AU Metzler, David E.
TI trace minerals, chelating agents, and the porphyrias
AU Peters, Henry A.
IT 1499-45-2 13933-92-1 13933-97-6
13934-03-7 57212-58-5 63221-70-5
91200-59-8 93353-85-6

L10 ANSWER 10 OF 16 CAOLD COPYRIGHT 2008 ACS on STN
AN CA53:7165b CAOLD
TI furoyl and furfuyl derivs. of pyridoxamine
AU McCasland, G. E.; Blanz, E., Jr.; Furst, A.
IT 4664-26-0 102313-26-8 103649-84-9 109401-44-7 114133-79-8
114493-09-3

L10 ANSWER 11 OF 16 CAOLD COPYRIGHT 2008 ACS on STN
AN CA52:2960g CAOLD
TI protective effect of N-pyridoxylidene-L-cysteine against x-ray irradiation
AU Yamada, Kozo; Hayami, S.; Sawaki, S.
IT 13933-88-5

L10 ANSWER 12 OF 16 CAOLD COPYRIGHT 2008 ACS on STN
AN CA51:18006i CAOLD
TI 4-pyridoxylamino-3-isoxazolidinones
PA Merck & Co., Inc.
DT Patent
TI 4-pyridoxylamino-3-isoxazolidones
AU Folkers, Karl
DT Patent
PATENT NO. KIND DATE

PI US 2801248 1957
IT 101495-73-2 101568-92-7 101655-14-5
106273-77-2

L10 ANSWER 13 OF 16 CAOLD COPYRIGHT 2008 ACS on STN
AN CA51:8804h CAOLD
TI 4-pyridoxylamino-3-isoxazolidinones
PA Merck & Co., Inc.
DT Patent
TI 4-pyridoxylamino-3-isoxazolidones
AU Folkers, Karl
DT Patent
PATENT NO. KIND DATE

PI US 2776296 1957
IT 101495-73-2 101568-92-7 101655-14-5 102015-45-2
106273-77-2

L10 ANSWER 14 OF 16 CAOLD COPYRIGHT 2008 ACS on STN
AN CA51:5870i CAOLD
TI equilibrium between pyridoxal and amino acids and their imines
AU Metzler, David E.
IT 1499-45-2 6956-94-1 7146-98-7
13933-86-3 13933-92-1 13933-97-6
13934-01-5 13934-03-7 17390-01-1
19973-35-4 57212-58-5 57237-43-1

Updated Search

10551430

63221-70-5 74317-99-0 91200-59-8
91761-12-5 93353-85-6 93688-50-7
100377-38-6 102015-20-3

L10 ANSWER 15 OF 16 CAOLD COPYRIGHT 2008 ACS on STN
AN CA51:587i CAOLD
TI biochem. aspects of atherosclerosis
AU Anfinsen, Christian B.
IT 57211-84-4

L10 ANSWER 16 OF 16 CAOLD COPYRIGHT 2008 ACS on STN
AN CA51:587b CAOLD
TI acute nephrosis following bleeding caused by lack of fibrin
AU Runge, Hans; Pfau, P.
IT 57211-84-4

=> FIL REGISTRY

COST IN U.S. DOLLARS	SINCE FILE ENTRY	TOTAL SESSION
FULL ESTIMATED COST	11.50	328.03

DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS)	SINCE FILE ENTRY	TOTAL SESSION
CA SUBSCRIBER PRICE	0.00	-8.00

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STRUCTURE FILE UPDATES: 19 AUG 2008 HIGHEST RN 1042061-07-3
DICTIONARY FILE UPDATES: 19 AUG 2008 HIGHEST RN 1042061-07-3

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http://www.cas.org/support/stngen/stndoc/properties.html

=> S 57211-84-4/RN

L11 1 57211-84-4/RN

Updated Search

10551430

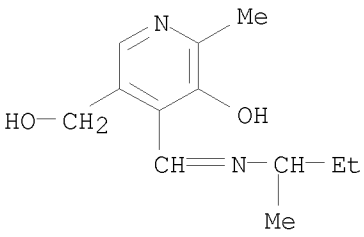
=> SET NOTICE 1 DISPLAY

NOTICE SET TO 1 U.S. DOLLAR FOR DISPLAY COMMAND
SET COMMAND COMPLETED

=> D L11 SQIDE 1-

YOU HAVE REQUESTED DATA FROM 1 ANSWERS - CONTINUE? Y/(N):y
THE ESTIMATED COST FOR THIS REQUEST IS 6.65 U.S. DOLLARS
DO YOU WANT TO CONTINUE WITH THIS REQUEST? (Y)/N:y

L11 ANSWER 1 OF 1 REGISTRY COPYRIGHT 2008 ACS on STN
RN 57211-84-4 REGISTRY
CN 3-Pyridinemethanol, 5-hydroxy-6-methyl-4-[[(1-methylpropyl)imino]methyl]-
(CA INDEX NAME)
OTHER CA INDEX NAMES:
CN 3-Pyridinemethanol, 4-(N-sec-butylformimidoyl)-5-hydroxy-6-methyl- (6CI)
MF C12 H18 N2 O2
LC STN Files: BEILSTEIN*, CA, CAOLD, CAPLUS
(*File contains numerically searchable property data)
DT.CA CAplus document type: Journal
RL.NP Roles from non-patents: PREP (Preparation); PRP (Properties); RACT
(Reactant or reagent)



PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

3 REFERENCES IN FILE CA (1907 TO DATE)
3 REFERENCES IN FILE CAPLUS (1907 TO DATE)
2 REFERENCES IN FILE CAOLD (PRIOR TO 1967)

=> SET NOTICE LOGIN DISPLAY

NOTICE SET TO OFF FOR DISPLAY COMMAND
SET COMMAND COMPLETED

Updated Search